immediate inspection at the Federal Reserve Bank(s) indicated below and at the offices of the Board of Governors. This information may also be obtained on an expedited basis, upon request, by contacting the appropriate Federal Reserve Bank and from the Board's Freedom of Information Office at *https://www.federalreserve.gov/foia/request.htm.* Interested persons may express their views in writing on the standards enumerated in the BHC Act (12 U.S.C. 1842(c)).

Comments regarding each of these applications must be received at the Reserve Bank indicated or the offices of the Board of Governors, Ann E. Misback, Secretary of the Board, 20th Street and Constitution Avenue NW, Washington, DC 20551–0001, not later than May 4, 2022.

A. Federal Reserve Bank of Philadelphia (William Spaniel, Senior Vice President) 100 North 6th Street, Philadelphia, Pennsylvania 19105–1521. Comments can also be sent electronically to *Comments.applications@phil.frb.org:*

1. *Fulton Financial Corporation, Lancaster, Pennsylvania;* to merge with Prudential Bancorp, Inc., and thereby indirectly acquire Prudential Bank, both of Philadelphia, Pennsylvania.

Board of Governors of the Federal Reserve System, March 30, 2022.

Michele Taylor Fennell,

Deputy Associate Secretary of the Board.

[FR Doc. 2022–07057 Filed 4–1–22; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000–0082; Docket No. 2022–0053; Sequence No. 11]

Information Collection; Federal Acquisition Regulation Part 7 Requirements

AGENCY: Department of Defense (DOD), General Services Administration (GSA), and National Aeronautics and Space Administration (NASA).

ACTION: Notice and request for comments.

SUMMARY: In accordance with the Paperwork Reduction Act of 1995, and the Office of Management and Budget (OMB) regulations, DoD, GSA, and NASA invite the public to comment on a revision concerning Federal Acquisition Regulation part 7 requirements. DoD, GSA, and NASA invite comments on: Whether the proposed collection of information is necessary for the proper performance of the functions of Federal Government acquisitions, including whether the information will have practical utility; the accuracy of the estimate of the burden of the proposed information collection; ways to enhance the quality, utility, and clarity of the information to be collected; and ways to minimize the burden of the information collection on respondents, including the use of automated collection techniques or other forms of information technology. OMB has approved this information collection for use through August 31, 2022. DoD, GSA, and NASA propose that OMB extend its approval for use for three additional years beyond the current expiration date.

DATES: DoD, GSA, and NASA will consider all comments received by June 3, 2022.

ADDRESSES: DoD, GSA, and NASA invite interested persons to submit comments on this collection through *https://www.regulations.gov* and follow the instructions on the site. This website provides the ability to type short comments directly into the comment field or attach a file for lengthier comments. If there are difficulties submitting comments, contact the GSA Regulatory Secretariat Division at 202–501–4755 or *GSARegSec@gsa.gov.*

Instructions: All items submitted must cite OMB Control No. 9000–0082, Federal Acquisition Regulation Part 7 Requirements. Comments received generally will be posted without change to *https://www.regulations.gov,* including any personal and/or business confidential information provided. To confirm receipt of your comment(s), please check *www.regulations.gov,* approximately two-to-three days after submission to verify posting.

FOR FURTHER INFORMATION CONTACT: Carrie Moore, Procurement Analyst, at telephone 571–300–5917, or *carrie.moore@gsa.gov.*

SUPPLEMENTARY INFORMATION:

A. OMB Control Number, Title, and Any Associated Form(s)

9000–0082, Federal Acquisition Regulation Part 7 Requirements.

B. Need and Uses

DoD, GSA, and NASA are combining OMB Control Nos. for the Federal Acquisition Regulation (FAR) by FAR part. This consolidation is expected to improve industry's ability to easily and efficiently identify burdens associated with a given FAR part. The review of the information collections by FAR part allows improved oversight to ensure there is no redundant or unaccounted for burden placed on industry. Lastly, combining information collections in a given FAR part is also expected to reduce the administrative burden associated with processing multiple information collections.

This justification supports the revision of OMB Control No. 9000–0082 and combines it with the previously approved information collection under OMB Control No. 9000–0114, with the new title ''Federal Acquisition Regulation Part 7 Requirements''. Upon approval of this consolidated information collection, OMB Control No. 9000–0114 will be discontinued. The burden requirements previously approved under the discontinued number will be covered under OMB Control No. 9000–0082.

This clearance covers the information that offerors or contractors must submit to comply with the following FAR requirements:

FAR clause 52.207–3, Right of First Refusal of Employment, requires contractors to provide the contracting officer, within 120 days of beginning contract performance, the names of personnel who were: Adversely affected or separated from Government employment as a result of the contract award; and subsequently hired by the contractor to perform under the contract within 90 days after contract performance began. The information provided under this clause is used by the Government to ensure: Contractor compliance with providing the right of first refusal to such affected personnel; and certain obligations to displaced employees are met by the Government.

FAR provision 52.207–4, Economic Purchase Quantity—Supplies, permits offerors, who believe that acquisition of supplies in quantity different from what is being solicited would be more advantageous to the Government, to recommend with their offer a more economic purchase quantity for the required supplies. The information provided under this provision is used by the Government to acquire supplies at the total and unit costs most advantageous to the Government and to develop a database for future acquisitions of such items of supply.

C. Annual Burden

Respondents: 14,510.
Total Annual Responses: 14,510.
Total Burden Hours: 14,530.
Obtaining Copies: Requesters may obtain a copy of the information collection documents from the GSA

Regulatory Secretariat Division, by calling 202–501–4755 or emailing *GSARegSec@gsa.gov.* Please cite OMB Control No. 9000–0082, Federal Acquisition Regulation Part 7 Requirements.

Janet Fry,

Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

[FR Doc. 2022–06992 Filed 4–1–22; 8:45 am]

BILLING CODE 6820–EP–P

GOVERNMENT ACCOUNTABILITY OFFICE

Request for Nominations for the Board of Governors of the Patient-Centered Outcomes Research Institute (PCORI)

AGENCY: Government Accountability Office (GAO).

ACTION: Request for letters of nomination and resumes.

SUMMARY: The Patient Protection and Affordable Care Act gave the Comptroller General of the United States responsibility for appointing up to 21 members to the Board of Governors of the Patient-Centered Outcomes Research Institute. In addition, the Directors of the Agency for Healthcare Research and Quality and the National Institutes of Health, or their designees, are members of the Board. As the result of terms ending in September 2022, GAO is accepting nominations in the following categories: A surgeon, a state-licensed integrative health care practitioner, a representative of patients and health care consumers, a representative of device manufacturers or developers, a representative of pharmaceutical manufacturers or developers, and a representative of private payers who represents health insurance issuers. Nominations should be sent to the email address listed below. Acknowledgement of submissions will be provided within a week of submission.

DATES: Letters of nomination and resumes should be submitted no later than May 10, 2022, to ensure adequate opportunity for review and consideration of nominees prior to appointment.

ADDRESSES: Submit letters of nomination and resumes to *PCORI@gao.gov.* Include PCORI Nomination in the subject line of the email.

FOR FURTHER INFORMATION CONTACT: Ray Sendejas at (202) 512–7113 or *SendejasR@gao.gov* if you do not receive an acknowledgement or need additional information. For general information, contact GAO's Office of Public Affairs, (202) 512–4800.

Authority: Sec. 6301 and Sec. 10602, Pub. L. 111–148, 124 Stat. 119, 727, 1005 (2010); Div. N, Sec. 104, Pub. L. 116–94, 133 Stat. 2534 (2019).

Gene L. Dodaro,

Comptroller General of the United States.

[FR Doc. 2022–06452 Filed 4–1–22; 8:45 am]

BILLING CODE 1610–02–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket No. CDC–2022–0044]

CDC Recommendations for Hepatitis B Screening and Testing—United States, 2022; Request for Comment

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), located within the Department of Health and Human Services (HHS), announces the opening of a docket to obtain comment on proposed updated recommendations for hepatitis B virus (HBV) infection screening and testing (Proposed Updated Recommendations), including hepatitis B screening at least once in a lifetime for persons 18 years of age and older, using a three-test panel. The Proposed Updated Recommendations also expand existing risk-based testing recommendations to include the following populations, activities, exposures, or conditions associated with increased risk for HBV infection: Persons currently or formerly incarcerated in a jail, prison, or other detention setting; persons with a history of sexually transmitted infections or multiple sex partners; and persons with a history of hepatitis C virus infection. The Proposed Updated Recommendations are intended to inform the practices of and care by U.S. healthcare providers and are based on scientific evidence of the effectiveness and economic value of screening to diagnose current HBV infection among adults in the United States.

DATES: Written comments must be received on or before June 3, 2022.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2022–0044, by either of the following methods:

- *Federal eRulemaking Portal: http://www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Division of Viral Hepatitis, Centers for Disease Control and Prevention, 1600 Clifton Road NE, Mailstop U12–3, Atlanta, GA 30329, Attn: Docket No. CDC–2022–0044.

Instructions: All submissions received must include the agency name and Docket Number. All relevant comments received will be posted without change to *http://regulations.gov,* including any personal information provided. Do not submit comments by email; CDC does not accept comments by email. For access to the docket to read background documents or comments received, go to *http://www.regulations.gov.*

FOR FURTHER INFORMATION CONTACT: Erin Conners, Centers for Disease Control and Prevention, 1600 Clifton Road NE, Mailstop U12–3, Atlanta, GA 30329; Telephone: 404–639–8000; Email: *DVHpolicy@cdc.gov.*

SUPPLEMENTARY INFORMATION:

Public Participation

Interested persons or organizations are invited to participate by submitting written views, recommendations, and data related to any of the Proposed Updated Recommendations or supporting evidence. In addition, CDC invites comments specifically on the following questions:

- Based on the evidence presented in the full recommendations document (see the Supporting and Related Materials tab in the docket), does the evidence support the Proposed Updated Recommendations for HBV infection screening and testing? If not, please state the reason why and, if available, provide additional evidence for consideration.
- Are CDC's Proposed Updated Recommendations (see Supporting and Related Materials) clearly written? If not, what changes do you propose to make them clearer?
- If implemented as currently drafted, do you believe the Proposed Updated Recommendations would result in a reduction in HBV infections and associated health and financial consequences (*e.g.,* patient and healthcare costs to treat chronic hepatitis B) in the United States? If not, please provide an explanation and supporting data or evidence.

Please note that comments received, including attachments and other supporting materials, are part of the public record and are subject to public disclosure. Comments will be posted on *https://www.regulations.gov.* Therefore, do not include any information in your